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Patent US 201C1  
Attorney Docket No. 892,280-137

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:  
  
**CAVALERI et al.**  
  
Serial No. 10/828,379  
  
Filed: April 16, 2004  
  
For: METHODS OF ADMINISTERING  
DALBAVANCIN FOR  
TREATMENT OF BACTERIAL  
INFECTIONS

Group Art Unit: 1614

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450

Sir:

Pursuant to 37 C.F.R. §1.56 and in accordance with 37 C.F.R. §§1.97–1.98,  
information relating to the above–identified application is hereby disclosed. The  
accompanying Form PTO–1449 provides a listing of documents that may be relevant to the  
subject application.

It is requested that the Examiner fully consider the art cited in the accompanying Form  
1449, initial the left–most column of the form adjacent each cited reference, and return a copy

CERTIFICATE OF MAILING (37 C.F.R. §1.8a)

I hereby certify that this paper (along with any referred to as being attached or enclosed) is being deposited with the United States Postal Service on the date shown below with sufficient postage as First Class Mail in an envelope addressed to the Commissioner for Patents, P.O. Box 1450 Alexandria, VA 22313-1450.

September 17, 2004  
Date of Deposit  
IRI:1058069.1

*Cynthia B. Pacheco*  
Cynthia B. Pacheco

for Applicants' records. It is further requested that the art be cited on the cover of any patent issuing from the subject application..

This IDS is believed to be timely in that it is being submitted under 37 CFR § 1.97(b), that is (1) within three months of the filing date of the application, which is not a continued prosecution application filed under § 1.53(d); or (2) within three months of entry of the national stage as set forth in 37 CFR § 1.491; or (3) before the mailing of a first Office action on the merits; or (4) before the mailing of a first Office action after filing a request for continued examination under § 1.114. Thus, no fee is required..

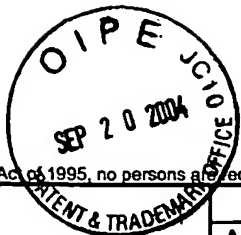
This statement should not be construed as a representation that more material information does not exist or that an exhaustive search of the relevant art has been made. Nor does this statement constitute an admission by Applicants or Applicants' agent that the information provided herein is necessarily prior art to Applicants' invention. Moreover, Applicants reserve the right to establish the patentability of the claimed invention over any of the listed documents should they be applied there-against as references..

Respectfully submitted,

O'MELVENY & MYERS LLP.

Dated: 9/17/04 By: Diane K. Wong  
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**INFORMATION DISCLOSURE  
STATEMENT BY APPLICANT**

(Use as many sheets as necessary)

**Complete if Known**

|                        |                  |
|------------------------|------------------|
| Application Number     | 10/828,379       |
| Filing Date            | April 16, 2004   |
| First Named Inventor   | CAVALERI et al.  |
| Art Unit               | 1614             |
| Examiner Name          | Not Yet Assigned |
| Attorney Docket Number | 892,280-137      |

Sheet 1 of 1

**NON PATENT LITERATURE DOCUMENTS**

| Examiner Initials* | Cite No. <sup>1</sup> | Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.                                 | T <sup>2</sup> |
|--------------------|-----------------------|---|----------------|
|                    | 1                     | Dowell, et al. (2003). "Dalbavancin Dosage Adjustments Not Required for Patients with Mild Renal Impairment," 2003 ECCMID Meeting.  |                |
|                    | 2                     | Stogniew et al. (2003). "Pharmacokinetic Attributes of Dalbavancin: Well Distributed and Completely Eliminated With Dual Routes of Elimination," 2003 ECCMID Meeting.   |                |
|                    | 3                     | White et al. (2000). "V-Glycopeptide: Phase1 Single and Multiple-dose Placebo Controlled Intravenous Safety, Pharmacokinetic, and Pharmacodynamic Study in Healthy Subjects," Abstracts of the 40th Interscience Conference on Antimicrobial Agents and Chemotherapy, September 2000, page 233. |                |
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| Examiner Signature | Date Considered |
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<sup>1</sup> Applicant's unique citation designation number (optional). <sup>2</sup> Applicant is to place a check mark here if English language Translation is attached.

This collection of information is required by 37 CFR 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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